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AMENDMENTS TO THE CLAIMS

1. (currently amended) A microarray hybridization device which comprises:

a flat substrate having a surface to which a microarray of reactive moieties

can be attached,

liquid barrier means juxtaposed with said surface to create a chamber

having an interior wall surface in which chamber said microarray is located, and

a cover means closing said chamber so said device may be manipulated

without loss of liquid target solution that fills said chamber except for a gaseous bubble

included therein,

said barrier means having inwardly facing surfaces which border said

chamber and which are generally perpendicular thereto, which surfaces are formed with a

plurality of bubble-fracturing elements that lie in a planar region extending parallel to

said flat substrate and extend laterally into said chamber so that, when said device is

moved so that a liquid target solution in said chamber moves along said flat substrate

surface from one boundary of said chamber to another boundary, a bubble initially in said

chamber is ruptured into a plurality of microbubbles that then assure very effective

distribution of the liquid target solution in said chamber across the entire microarray,

driven by movement of said microbubbles.

2. (original) The microarray hybridization device of claim 1 wherein said

cover is flat and is spaced uniformly from said surface by said barrier means.

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3. (original) The microarray hybridization device of claim 2 wherein said

cover is made of substantially rigid, transparent material.

4. (currently amended) The microarray hybridization device of claim 2

wherein said barrier means has a height such as to space said cover between about 0.2

and about 2 mm from said surface.

(currently amended) The microarray hybridization device of claim 2

wherein said barrier means forms a generally rectangular perimeter of said chamber

having four walls and wherein one or more of the four walls of said barrier means

includes sharp edges that are aligned substantially perpendicular to said surface upon

which the microarray is attached, which edges are spaced apart by pockets and function

as said bubble-fracturing elements.

(original) The microarray hybridization device according to claim 5

wherein said bubble-fracturing elements are disposed along two opposed boundary walls

of said rectangular perimeter barrier and are formed by a plurality of generally triangular

fingers that project from boundary walls into said chamber and have said sharp edges at

the tips thereof, with said pockets being located therebetween.

7. (currently amended) The microarray hybridization device according to

claim 5 $\underline{6}$ wherein said rectangular perimeter includes two longer walls and two shorter

walls with said bubble-fracturing elements being formed as part of said two shorter walls.

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8. (original) The microarray hybridization device according to claim 7 wherein said

triangular fingers in said two shorter walls are aligned so as to project in the direction

from which bubbles in the target solution in said chamber will normally approach the

respective wall when the device moved during hybridization.

(original) The microarray hybridization device according to claim 2

wherein said bubble-fracturing elements are formed of hydrophobic material.

10. (original) The microarray hybridization device according to claim 2

wherein said cover is made of an opaque hydrophobic material and includes at least one

filling port through which said liquid target solution can be supplied into said chamber

wherein a microarray is disposed.

11. (currently amended) A microarray hybridization device which comprises:

a flat substrate having an upper surface,

a microarray of reactive moieties attached to said upper surface,

a liquid perimeter barrier juxtaposed with said surface to create a chamber

having an interior wall surface, in which chamber said microarray is located, and

a cover juxtaposed with said barrier to close said chamber so said device

may be manipulated without loss of a liquid target solution that fills said chamber except

for a gaseous bubble included therein,

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said perimeter barrier having inwardly facing walls which border said

chamber and which are generally perpendicular thereto, which walls are formed with a

plurality of bubble fracturing elements that lie in a planar region extending parallel to

said flat substrate and extend laterally into said chamber so that, when said device is

moved so that the liquid target solution moves along said flat substrate surface upon

which said microarray is located, a bubble initially in said chamber is ruptured into a

plurality of microbubbles that assure very effective distribution of a the liquid target

solution in said chamber across the entire microarray, driven by movement of said

microbubbles.

12. (original) The microarray hybridization device of claim 11 wherein said

cover is flat, being made of substantially rigid, transparent material, and is spaced

uniformly about 0.2 and about 2 mm from said surface by said perimeter barrier.

13. (original) The microarray hybridization device of claim 12 wherein said

perimeter barrier forms a generally rectangular chamber and wherein one or more of the

four walls thereof includes protrusions having sharp edges that are aligned substantially

perpendicular to said surface on which said microarray is located, said protrusions being

spaced apart by pockets and functioning as said bubble-fracturing elements.

14. (original) The microarray hybridization device according to claim 12

wherein said cover includes at least one filling port through which said liquid target

solution can be supplied into said chamber and wherein said microarray includes a

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plurality of 3D spots which are attached to said upper surface and extend upward therefrom at least about 20 µm, which 3D spots carry said reactive moieties.

15-17. (canceled).

18. (withdrawn-currently amended) A method of effecting hybridization between probes and a target solution, which method comprises:

between process and a tanget botation, which method comprises

hybridization device according to claim 1 containing a microarray of reactive probe

providing a flat substrate having a surface to which microarray

moieties are attached,

juxtaposing a perimeter liquid barrier with said surface to create a chamber, in which said microarray is located, and closing said chamber so said substrate may be manipulated without loss of liquid target solution,

filling said chamber of said device with a target solution and a gaseous bubble, and

moving said substrate to cause the target solution to move from one boundary of said chamber to another with at least one such boundary being shaped so that as a result of such movement the bubble in said chamber is ruptured into a plurality of microbubbles that then assure very effective distribution of the liquid target solution across the entire microarray, driven by subsequent movement of such microbubbles.

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19. (withdrawn) The method of claim 18 wherein said chamber is formed and

closed by a flat cover and a depending perimeter gasket that spaces said cover uniformly

from said surface.

20. (withdrawn) The method of claim 19 wherein said gasket forms a

generally rectangular perimeter of said chamber having two opposed shorter walls and

wherein said microbubbles are created by sharp-edged protrusions that project into said

chamber from said two opposed shorter walls in the direction from which a bubble would

approach each said wall during normal movement.

21. (withdrawn) The method according to claim 20 wherein said target

solution is introduced through at least one filling port in said cover which is then sealed.

22. (withdrawn) The method of claim 20 wherein said substrate is moved by

rotation about an axis which is substantially horizontal and wherein said chamber is

aligned so that said shorter walls are generally perpendicular to a line extending radially

from said axis of rotation.

23. (new) A microarray hybridization device which comprises:

a flat substrate having an upper surface,

a microarray of reactive moieties in 3D spots attached to said upper

surface.

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a rectangular liquid perimeter barrier juxtaposed with said surface to

create a chamber in which said microarray is located,

a flat cover juxtaposed with said barrier to close said chamber so as to

confine a liquid target solution therein that fills said chamber except for a gaseous

bubble,

said cover including at least one filling port through which a liquid target

solution can be supplied into said chamber, and

means for sealing said port so said device may be manipulated in a

substantially vertical plane about a horizontal axis without loss of the confined target

solution,

said perimeter barrier having four inwardly facing surfaces which border

said chamber and which are generally perpendicular thereto, which surfaces are formed

with a plurality of bubble-fracturing elements that extend laterally into said chamber in a

planar region between said cover and said flat substrate, so that, when said device is

manipulated causing the liquid target solution to move along said flat substrate surface

upon which said microarray is located, a bubble initially in said chamber is ruptured into

a plurality of microbubbles that assure very effective distribution of the liquid target

solution in said chamber across the entire microarray, driven by movement of said

microbubbles.

24. (new) The microarray hybridization device of claim 23 wherein said

bubble-fracturing elements comprise generally triangular protrusions having sharp edges,

that are aligned substantially perpendicular to said surface on which said microarray is

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located, which elements are located on one or more of four walls which form said perimeter barrier, said generally triangular protrusions being spaced apart by pockets.

25. (new) The microarray hybridization device according to claim 24 wherein said protrusions are located on two shorter walls of said four walls of said rectangular barrier and wherein said plurality of 3D spots which are attached to said upper surface extend upward therefrom at least about 20 μ m.